On January 13, 1942, the United States attorney for the District of Oregon filed a libel against 115 ¼-ounce packages of Wemett's Salve at Portland, Oreg., alleging that the article had been shipped on or about August 28 and October 1, 1941, by F. J. Wemett from Los Angeles, Calif.; and charging that it was misbranded in that its container was so made, formed, and filled as to be misleading.

On March 25, 1942, no claimant having appeared, judgment of condemnation was entered and the product was ordered destroyed, or delivered to a charitable institution.

NONSTERILE SURGICAL DRESSINGS

698. Adulteration and misbranding of sutures. U. S. v. 32 Packages of Sutures. Default decree of condemnation and destruction. (F. D. C. No. 6762. Sample No. 71511-E.)

On January 26, 1942, the United States attorney for the Southern District of Iowa filed a libel against the above-named product at Des Moines, Iowa, alleging that it had been shipped on or about September 17, 1941, by Davis Sutures, Inc., from Chicago, Ill.; and charging that it was adulterated and misbranded.

The article was alleged to be adulterated in that it purported to be and was represented as a drug the name of which is recognized in the United States Pharmacopoeia, but its purity fell below the standard set forth in the pharmacopoeia since it was not sterile. It was alleged to be misbranded in that the statement in the labeling, "Guaranty Davis Sutures are guaranteed to be sterile," was false and misleading since it was not sterile but was contaminated with viable aerobic and anaerobic or facultative anaerobic micro-organisms, including spore-bearing and gas-producing micro-organisms.

On February 28, 1942, no claimant having appeared, judgment of condemnation

was entered and the product was ordered destroyed.

699. Misbranding of finger compresses. U. S. v. 1,344 Packages of Quick Strips Finger Compresses. Default decree of condemnation and destruction. (F. D. C. No. 6901. Sample Nos. 92009–E, 92010–E.)

On February 20, 1942, the United States attorney for the Southern District of California filed a libel against the above-named product at Los Angeles, Calif., alleging that it had been shipped on or about January 23, 1942, by the Quick Manufacturing Co. from Chicago, Ill.; and charging that it was misbranded.

The article was alleged to be misbranded in that designs showing application of the strips to the finger and the statements, "Place Medicated Pad over Injury," "Press Edges Together," "Wrap Around Finger," and "Medicated With Boric Acid or Iodochrome," were misleading when applied to a bandage which was contaminated with viable micro-organisms; and in that such designs and statements suggested that it would be suitable for first aid purposes; whereas it was not.

On March 19, 1942, no claimant having appeared, judgment of condemnation was entered and the product was ordered destroyed.

700. Adulteration and misbranding of Hill's Swabbed Applicators with Tongue Blade. U. S. v. 76 Cartons of Hill's Swabbed Applicators with Tongue Blade. (F. D. C. No. 6849. Sample No. 70098–E.)

On or about March 2, 1942, the United States attorney for the Southern District of Florida filed a libel against 76 cartons of the above-named product at Jacksonville, Fla., alleging that it had been shipped on or about November 27, 1941, by the Wetmore-Century Corporation from New York, N. Y.; and charging that it was adulterated and misbranded.

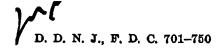
The article was alleged to be adulterated in that its purity and quality fell below that which it purported or was represented to possess, namely, (glassine envelope) "sterilized," since it was not sterile but was contaminated with aerobic, anaerobic, or facultative anaerobic micro-organisms.

It was alleged to be misbranded in that the following statements in the labeling, (envelope) "Sterilized Applicators * * * Sterilized After Packing," and (carton) "The Modern Way of Treating sore throats, cuts, wounds, ear and nose ailments. The Ideal Way of safeguarding your health * * * For eye, ear and nose treatment * * * especially useful to mothers treating infants * * * specially made for Throat Treatment," were false and misleading when applied to an article that was not sterile but was contaminated with viable micro-organisms.

On March 21, 1942, no claimant having appeared, judgment of condemnation was entered and the product was ordered destroyed.

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FEDERAL SECURITY AGENCY

FOOD AND DRUG ADMINISTRATION

NOTICES OF JUDGMENT UNDER THE FEDERAL FOOD, DRUG, AND COSMETIC ACT

[Given pursuant to section 705 of the Food, Drug, and Cosmetic Act]

701-750

DRUGS AND DEVICES

The cases reported herewith were instituted in the United States district courts by the United States attorneys acting upon reports submitted by direction of the Federal Security Administrator.

WATSON B. MILLER, Acting Administrator, Federal Security Agency. WASHINGTON, D. C., February 12, 1943.

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DRUGS ACTIONABLE BECAUSE OF POTENTIAL DANGER WHEN USED ACCORDING TO DIRECTIONS

701. Action to restrain interstate shipment of Alcoban, a misbranded drug. U. S. v. Maffett Sales Corporation, Frank L. Wilson, Nell B. Wilson, and Reuel K. Yount. Temporary restraining order entered. Default order granting permanent injunction. (Inj. No. 17.)

On October 20, 1941, the United States attorney for the Western District of Washington filed a complaint against the Maffet Sales Corporation and Frank L. Wilson, Nell B. Wilson, and Reuel K. Yount, Seattle, Wash., alleging that the defendants for many years past, had been engaged in the sale and distribution of an article of drugs called Alcoban; that the article was sold by the defendants in cartons which bore the printed statement, "An Aid in Curbing the Liquor Habit," and was accompanied by a circular which contained, among others, the representation that it was an aid in curing the liquor habit, and directions that the contents of 1 capsule should be given every 15 to 20 minutes until 3 capsules were taken; that, if vomiting occurred, this should be regarded as a proper dosage; that, if no vomiting occurred on the 1-capsule per drink basis, the dosage should be doubled, and if vomiting then occurred this should be considered the correct dosage; and that, if no vomiting occurred after the consumption of three single-dose drinks and two double-dose drinks, the treatment should be discontinued. The complaint alleged further that the statements in the labeling were false and misleading since the article did not constitute an appropriate remedy for the purposes stated and recommended; that the use and administration of the drug in the dosage and with the frequency and duration prescribed, recommended, and suggested in the labeling was dangerous to health, and that consequently the product was misbranded. The complaint alleged further that

[D. D. N.]

the defendants at that time were introducing and delivering the said drug for introduction into interstate commerce and prayed that judgment and decree be entered permanently restraining and enjoining them and all acting upon their behalf from continuing to do so; and prayed that a preliminary injunction be granted restraining the defendants during the pendency of the action.

On November 10, 1941, the court granted a temporary restraining order in accordance with the prayer of the complaint. On June 9, 1942, the defendants then being in default, judgment was entered permanently and forever enjoining and restraining them from directly or indirectly introducing or delivering for

introduction said drug into interstate commerce.

702. Misbranding of Lambert's Powders. U. S. v. Claude M. Stanley (Stanley Drug Co.). Plea of guilty. Fine, \$50. (F. D. C. No. 4161. Sample No. 38881-E.)

This product when used according to directions on the label, would be dangerous to health, the label failed to bear adequate warning statements, and it

also contained false and misleading claims.

On November 10, 1941, the United States attorney for the District of Minnesota filed an information against Claude M. Stanley, trading as the Stanley Drug Co. at Minneapolis, Minn., alleging shipment on or about July 19, 1940, from the State of Minnesota into the State of Wisconsin of a quantity of Lambert's Powders that were misbranded.

Analysis of a sample of the article showed that each powder contained

acetanilid (2½ grains), aspirin (5 grains), and salol (2½ grains).

The article was alleged to be misbranded: (1) In that it was dangerous to health when used in the dosage or with the frequency or duration prescribed, recommended, or suggested in the labeling, i. e., "Directions * * * Adult Dose: One before each meal and one at bedtime." (2) In that its labeling failed to bear adequate warnings against use by children where its use might be dangerous to health, or against unsafe dosage or methods or duration of administration in such manner and form as are necessary for the protection of users since each powder contained approximately 2½ grains of acetanilid, and the labeling did not bear a warning that frequent or continuous use might cause serious blood disturbances, anemia, collapse, or a dependence on the drug, and that it should not be given to children. (3) In that the statement (carton) "muscular aches and body pains, lumbago," was false and misleading since it represented that the drug was efficacious in the treatment of muscular aches, body pains, and lumbago; whereas it was not efficacious for such purposes.

On March 3, 1942, the defendant entered a plea of guilty and the court

imposed a fine of \$50.

703. Misbranding of a.m. Solution. U. S. v. 7½ Dozen Packages of a.m. Solution. Default decree of condemnation and destruction. (F. D. C. No. 6839. Sample No. 79171–E.)

This product contained chrysarobin and would be dangerous to health when used according to directions. Its label also contained false and misleading

therapeutic claims.

On February 13, 1942, the United States attorney for the Middle District of Tennessee filed a libel against the above-named product at Nashville, Tenn., alleging that it had been shipped on or about November 13, 1941, and January 14, 1942, by the Kenton Pharmacal Co., Inc., from Covington, Ky.; and charging that it was misbranded.

Analysis of a sample of the article showed that it consisted essentially of chrysarobin (approximately 0.66 grain per fluid ounce), salicylic acid, benzoic

acid, alcohol, and a volatile oil.

The article was alleged to be misbranded: (1) In that it was dangerous to health when used in the dosage or with the frequency or duration prescribed or recommended in the labeling. (2) In that the following statements, "For the relief of itching and discomfort of Athlete's Foot (Dermatophytosis), Ringworm, Insect Bites, Impetigo, externally caused Eczema, Rashes and Pimples, and other forms of local skin irritations," were false and misleading since they represented and suggested that when used as directed it constituted a safe and efficacious treatment for the relief of the itching torment and discomfort of athlete's foot and other skin irritations named above; whereas it was not safe when used as directed and was not an efficacious treatment for such conditions.

On April 9, 1942, no claimant having appeared, judgment of condemnation

was entered and the product was ordered destroyed.